



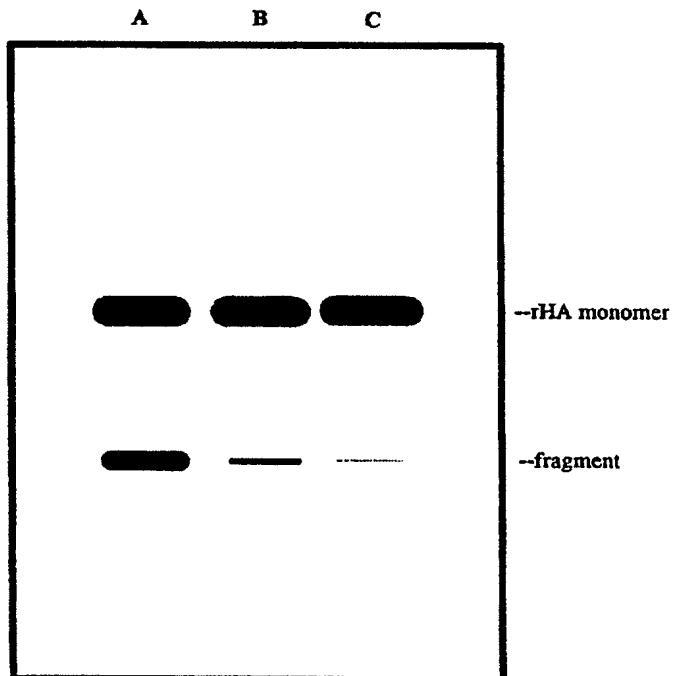
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(54) Title: YEAST STRAINS AND MODIFIED ALBUMINS

(57) Abstract

Albumin, for example human albumin, is expressed and secreted in yeast which has been mutated to lack the yeast aspartyl protease 3 (Yap3p) or its equivalent, thereby reducing the production of a 45kD albumin fragment. A further reduction is achieved by additionally deleting the Kex2p function. Alternatively, a modified albumin is prepared which is not susceptible to Yap3p cleavage, for example human albumin which is R410A, K413Q and K414Q.



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YEAST STRAINS AND MODIFIED ALBUMINS

Field of the invention

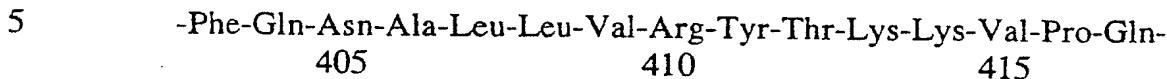
- 5 The present invention relates to the production of recombinant human albumin (rHA) by yeast species.

Background and prior art

- 10 Human serum albumin (HSA) is a protein of 585 amino acids that is responsible for a significant proportion of the osmotic pressure of serum, and also functions as a carrier of endogenous and exogenous ligands. It is used clinically in the treatment of patients with severe burns, shock, or blood loss, and at present is produced commercially by extraction from human blood. The
15 production of recombinant human albumin (rHA) in microorganisms has been disclosed in EP 330 451 and EP 361 991.

- In recent years yeast species have been widely used as a host organisms for the production of heterologous proteins (reviewed by Romanos *et al*, 1992),
20 including rHA (Sleep *et al*, 1990, 1991; Fleer *et al*, 1991). Yeasts are readily amenable to genetic manipulation, can be grown to high cell density on simple media, and as eukaryotes are suitable for production of secreted as well as cytosolic proteins.
- 25 When *S. cerevisiae* is utilised to produce rHA, the major secreted protein is mature 67kDa albumin. However, a 45kDa N-terminal fragment of rHA is also observed (Sleep *et al*, 1990). A similar fragment is obtained when rHA is expressed in *Kluyveromyces* sp. (Fleer *et al*, 1991) and *Pichia pastoris* (EP 510 693). The fragment has the same N-terminal amino acid sequence as
30 mature rHA, but the carboxy terminus is heterogeneous and occurs between

Phe⁴⁰³ and Val⁴⁰⁹ with the most common termini being Leu⁴⁰⁷ and Val⁴⁰⁹ (Geisow *et al*, 1991), as shown below.



- The amount of fragment produced, as a percentage of total rHA secreted,
10 varies with both the strain and the secretion leader sequence utilised, but is
never reduced to zero (Sleep *et al*, 1990). We have also found that the amount
of fragment produced in high cell density fermentation (75-100g/L cell dry
weight) is approximately five times higher than in shake flask cultures.
- 15 The 45kDa albumin fragment is not observed in serum-derived human serum
albumin (HSA), and its presence as non-nature-identical material in the
recombinant product is undesirable. The problem addressed by the present
invention is to reduce the amount of the 45kDa fragment in the product. The
simplest and most obvious approach would have been to have purified it away
20 from the full length albumin, as proposed by Gist-brocades in EP 524 681 (see
especially page 4, lines 17-22). However, we have chosen a different
approach, namely to try to avoid its production in the first place.
- Sleep *et al* (1990) postulated that rHA fragment is produced within the cell and
25 is not the result of extra-cellular proteolysis. These authors codon-optimised
the HSA cDNA from Glu³⁸² to Ser⁴¹⁹ but this had no effect on production of
rHA fragment. They noted that a potential Kex2p processing site in the rHA
amino acid sequence, Lys⁴¹³Lys⁴¹⁴, is in close proximity to the heterogeneous
carboxy terminus of the fragment, but neither use of a *kex2* host strain (ie a
30 strain harbouring a mutation in the *KEX2* gene such that it does not produce the
Kex2p protease), nor removal of the potential cleavage site by site-directed

mutagenesis of the codon for Lys⁴¹⁴, resulted in reduction in the amount of the fragment.

There is a vast array of yeast proteases which could, in principle, be degrading
5 a desired protein product, including (in *S. cerevisiae*) yscA, yscB, yscY, yscS,
other vacuolar proteinases, yscD, yscE, yscF (equivalent to kex2p), ysc α ,
yscIV, yscG, yscH, yscJ, yscE and kex1.

Bourbonnais *et al* (1991) described an *S. cerevisiae* endoprotease activity
10 specific for monobasic sites, an example of which (Arg⁴¹⁰) exists in this region
of albumin. This activity was later found to be attributable to yeast aspartyl
protease 3 (Yap3) (Bourbonnais *et al*, 1993), an enzyme which was originally
described by Egel-Mitani *et al* (1990) as an endoprotease similar to Kex2p in
specificity, in that it cleaved at paired basic residues. Further work suggested
15 that Yap3p is able to cleave monobasic sites and between, and C-terminal to,
pairs of basic residues, but that cleavage at both types of sites is dependent on
the sequence context (Azaryan *et al*, 1993; Cawley *et al*, 1993).

As already discussed, the region of the C-terminus of rHA fragment contains
20 both a monobasic (Arg⁴¹⁰) and a dibasic site (Lys⁴¹³Lys⁴¹⁴). However, even
though a Kex2p-like proteolytic activity is present in human cells and is
responsible for cleavage of the pro sequence of HSA C-terminal to a pair of
arginine residues, the fragment discussed above is not known to be produced
in humans. This indicates that the basic residues Arg⁴¹⁰, Lys⁴¹³ and Lys⁴¹⁴ are
25 not recognised by this Kex2p-like protease, in turn suggesting that this region
of the molecule may not be accessible to proteases in the secretory pathway.
Thus, the Yap3p protease could not have been predicted to be responsible for
the production of the 45kDa fragment. In addition, Egel-Mitani *et al* (1990
Yeast 6, 127-137) had shown Yap3p to be similar to Kex2p in cleaving the
30 MF α prohormone. Since removal of the Kex2p function alone does not

reduce the amount of the fragment produced, there was no reason to suppose that removal of the Yap3p function would be beneficial. Indeed, Bourbonnais *et al* (1993) showed that *yap3* strains had a decreased ability to process prosomatostatin, and therefore taught away from using *yap3* strains in the 5 production of heterologous proteins.

Summary of the invention

The solution to the problem identified above is, in accordance with the 10 invention, to avoid or at least reduce production of the fragment in the initial fermentation, rather than to remove it during purification of the albumin. We have now found that, out of the 20 or more yeast proteases which are so far known to exist, it is in fact the Yap3p protease which is largely responsible for the 45kD fragment of rHA produced in yeast. The present invention provides 15 a method for substantially reducing the amount of a 45kDa fragment produced when rHA is secreted from yeast species. The reduction in the amount of fragment both improves recovery of rHA during the purification process, and provides a higher quality of final product. A further, and completely unexpected, benefit of using *yap3* strains of yeast is that they can produce 30- 20 50% more rHA than strains having the Yap3p function. This benefit cannot be accounted for merely by the reduction of rHA fragment from ~ 15% to 3-5%.

Thus, one aspect of the present invention provides a process for preparing 25 albumin by secretion from a yeast genetically modified to produce and secrete the albumin, comprising culturing the yeast in a culture medium such that albumin is secreted into the medium, characterised in that the yeast cells have a reduced level of yeast aspartyl protease 3 proteolytic activity.

Preferably, the said proteolytic activity is an endoprotease activity specific for 30 monobasic sites and for paired basic amino acids in a polypeptide.

Suitably, the yeast is *S. cerevisiae* which lacks a functional *YAP3* gene. However, the invention is not limited to the use of *S. cerevisiae*, since the problem of 45 kDa fragment production is found also in other yeast genera, for example *Pichia* and *Kluyveromyces*, which shows that they have equivalent proteases (ie Yap3p proteolytic activity); see Clerc *et al* (1994), page 253. We have confirmed this by hybridisation analysis to locate homologues of Yap3p in non-*Saccharomyces* genera. A gene is regarded as a homologue, in general, if the sequence of the translation product has greater than 50% sequence identity to Yap3p. In non-*Saccharomyces* genera, the Yap3p-like protease and its gene may be named differently, but this does not of course alter their essential nature.

The level of fragment can be reduced still further if, as well as substantially eliminating the Yap3p proteolytic activity, the Kex2p function is also substantially eliminated even though, as mentioned above, elimination of the Kex2p function alone does not affect the level of fragment. As in the case of Yap3p, the Kex2p function is not restricted to *Saccharomyces*; see Gellissen *et al* (1992), especially the sentence bridging pages 415 and 416, showing that *Pichia* has a Kex2p function. The genes encoding the Kex2p equivalent activity in *Kluyveromyces lactis* and *Yarrowia lipolytica* have been cloned (Tanguy-Rougeau *et al*, 1988; Enderlin & Ogrydziak, 1994).

A suitable means of eliminating the activity of a protease is to disrupt the host gene encoding the protease, thereby generating a non-reverting strain missing all or part of the gene for the protease (Rothstein, 1983). Alternatively, the activity can be reduced or eliminated by classical mutagenesis procedures or by the introduction of specific point mutations by the process of transplacement (Winston *et al*, 1983). Preferably, the activity of the enzyme is reduced to at most 50% of the wild-type level, more preferably no more than 25%, 10% or 30%, and most preferably is undetectable. The level of Yap3p proteolytic

activity may be measured by determining the production of the 45 kDa fragment, or by the ^{125}I - β_h -lipoprotein assay of Azaryan *et al* (1993), also used by Cawley *et al* (1993). Kex2p proteolytic activity may similarly be measured by known assays, for example as set out in Fuller *et al* (1989).

5

The albumin may be a human albumin, or a variant thereof, or albumin from any other animal.

- By "variants" we include insertions, deletions and substitutions, either 10 conservative or non-conservative, where such changes do not substantially alter the oncotic, useful ligand-binding or non-immunogenic properties of albumin. In particular, we include naturally-occurring polymorphic variants of human 15 albumin; fragments of human albumin which include the region cleaved by Yap3p, for example those fragments disclosed in EP 322 094 (namely HSA (1-n), where n is 369 to 419) which are sufficiently long to include the Yap3p-cleaved region (ie where n is 403 to 419); and fusions of albumin (or Yap3p-cleavable portions thereof) with other proteins, for example the kind disclosed 20 in WO 90/13653.
- 25 By "conservative substitutions" is intended swaps within groups such as Gly, Ala; Val, Ile, Leu; Asp, Glu; Asn, Gln; Ser, Thr; Lys, Arg; and Phe, Tyr.

Such variants may be made using the methods of protein engineering and site-directed mutagenesis as described below.

25

- A second aspect of the invention provides a modified albumin having at least 90% sequence identity to a naturally-occurring albumin, which naturally-occurring albumin is susceptible to cleavage with the *S. cerevisiae* yeast 30 aspartyl protease 3 (Yap3p) when expressed in yeast, characterised in that the modified albumin is not susceptible to such cleavage.

Preferably, the modified albumin lacks a monobasic amino acid present in the naturally-occurring albumin protein. Suitably, the said monobasic amino acid is arginine. Conveniently, the modified albumin additionally lacks a pair of basic amino acids present in the naturally-occurring albumin, especially any of
5 Lys, Lys; Lys, Arg; Arg, Lys; or Arg, Arg. Thus, in one particular embodiment, the naturally-occurring albumin is human albumin and the modified protein lacks Arg⁴¹⁰ and, optionally, one or both Lys⁴¹³Lys⁴¹⁴ lysines. For example, the modified albumin may be human albumin having the amino acid changes R410A, K413Q, K414Q. Equivalent modifications in bovine
10 serum albumin include replacing the Arg⁴⁰⁸ and/or one or both of Arg⁴¹¹Lys⁴¹². The person skilled in the art will be able to identify monobasic sites and pairs of basic residues in other albumins without difficulty.

The numbering of the residues corresponds to the sequence of normal mature
15 human albumin. If the albumin is a variant (for example a polymorphic form) having a net deletion or addition of residues N-terminal to the position identified, then the numbering refers to the residues of the variant albumin which are aligned with the numbered positions of normal albumin when the two sequences are so aligned as to maximise the apparent homology.
20

A third aspect of the invention provides a polynucleotide encoding such a modified albumin.

The DNA is expressed in a suitable yeast (either the DNA being for a modified
25 albumin, or the yeast lacking the Yap3p function) to produce an albumin. Thus, the DNA encoding the albumin may be used in accordance with known techniques, appropriately modified in view of the teachings contained herein, to construct an expression vector, which is then used to transform an appropriate yeast cell for the expression and production of the albumin.

The DNA encoding the albumin may be joined to a wide variety of other DNA sequences for introduction into an appropriate host. The companion DNA will depend upon the nature of the host, the manner of the introduction of the DNA into the host, and whether episomal maintenance or integration is desired.

5

Generally, the DNA is inserted into an expression vector, such as a plasmid, in proper orientation and correct reading frame for expression. The vector is then introduced into the host through standard techniques and, generally, it will be necessary to select for transformed host cells.

10

Host cells that have been transformed by the recombinant DNA of the invention are then cultured for a sufficient time and under appropriate conditions known to those skilled in the art in view of the teachings disclosed herein to permit the expression and secretion of the albumin, which can then be recovered, as is known.

15

Useful yeast plasmid vectors are pRS403-406 and pRS413-416 and are generally available from Stratagene Cloning Systems, La Jolla, CA 92037, USA. Plasmids pRS403, pRS404, pRS405 and pRS406 are Yeast Integrating 20 plasmids (YIps) and incorporate the yeast selectable markers *HIS3*, *TRP1*, *LEU2* and *URA3*. Plasmids pRS413-416 are Yeast Centromere plasmids (YCps). Other yeast expression plasmids are disclosed in EP-A-258 067, EP-A-286 424 and EP-A-424 117.

25

The polynucleotide coding sequences encoding the modified albumin of the invention may have additional differences to those required to produce the modified albumin. For example, different codons can be substituted which code for the same amino acid(s) as the original codons. Alternatively, the substitute codons may code for a different amino acid that will not affect the 30 activity or immunogenicity of the albumin or which may improve its activity

or immunogenicity, as well as reducing its susceptibility to a Yap3p protease activity. For example, site-directed mutagenesis or other techniques can be employed to create single or multiple mutations, such as replacements, insertions, deletions, and transpositions, as described in Botstein and Shortle 5 (1985). Since such modified coding sequences can be obtained by the application of known techniques to the teachings contained herein, such modified coding sequences are within the scope of the claimed invention.

Exemplary genera of yeast contemplated to be useful in the practice of the 10 present invention are *Pichia*, *Saccharomyces*, *Kluyveromyces*, *Candida*, *Torulopsis*, *Hansenula* (now reclassified as *Pichia*), *Histoplasma*, *Schizosaccharomyces*, *Citeromyces*, *Pachysolen*, *Debaromyces*, *Metschunikowia*, *Rhodosporidium*, *Leucosporidium*, *Botryoascus*, *Sporidiobolus*, *Endomycopsis*, and the like. Preferred genera are those selected from the group consisting of 15 *Pichia*, *Saccharomyces*, *Kluyveromyces*, *Yarrowia* and *Hansenula*. Examples of *Saccharomyces* sp. are *S. cerevisiae*, *S. italicus* and *S. rouxii*. Examples of *Kluyveromyces* sp. are *K. fragilis* and *K. lactis*. Examples of *Hansenula* (*Pichia*) sp. are *H. polymorpha* (now *Pichia angusta*), *H. anomala* (now *P. anomala*) and *P. pastoris*. *Y. lipolytica* is an example of a suitable *Yarrowia* 20 species.

Methods for the transformation of *S. cerevisiae* are taught generally in EP 251 744, EP 258 067 and WO 90/01063, all of which are incorporated herein by reference. Suitable promoters for *S. cerevisiae* include those associated with 25 the *PGK1* gene, *GAL1* or *GAL10* genes, *CYC1*, *PHO5*, *TRP1*, *ADH1*, *ADH2*, the genes for glyceraldehyde-3-phosphate dehydrogenase, hexokinase, pyruvate decarboxylase, phosphofructokinase, triose phosphate isomerase, phosphoglucose isomerase, glucokinase, α -mating factor pheromone, α -mating factor pheromone, the *PRB1* promoter, the *GPD1* promoter, and hybrid 30 promoters involving hybrids of parts of 5' regulatory regions with parts of 5'

regulatory regions of other promoters or with upstream activation sites (eg the promoter of EP-A-258 067).

Convenient regulatable promoters for use in *Schizosaccharomyces pombe* are
5 the thiamine-repressible promoter from the *nmt* gene as described by Maundrell
(1990) and the glucose-repressible *fbp1* gene promoter as described by Hoffman
& Winston (1990).

Methods of transforming *Pichia* for expression of foreign genes are taught in,
10 for example, Cregg *et al* (1993), and various Phillips patents (eg US 4 857
467, incorporated herein by reference), and *Pichia* expression kits are
commercially available from Invitrogen BV, Leek, Netherlands, and Invitrogen
Corp., San Diego, California. Suitable promoters include *AOX1* and *AOX2*.

15 The Gellissen *et al* (1992) paper mentioned above and Gleeson *et al* (1986) *J.
Gen. Microbiol.* **132**, 3459-3465 include information on *Hansenula* vectors and
transformation, suitable promoters being *MOX1* and *FMD1*; whilst EP 361 991,
Fleer *et al* (1991) and other publications from Rhône-Poulenc Rorer teach how
to express foreign proteins in *Kluyveromyces* spp., a suitable promoter being
20 *PGK1*.

The transcription termination signal is preferably the 3' flanking sequence of
a eukaryotic gene which contains proper signals for transcription termination
and polyadenylation. Suitable 3' flanking sequences may, for example, be
25 those of the gene naturally linked to the expression control sequence used, ie
may correspond to the promoter. Alternatively, they may be different in which
case the termination signal of the *S. cerevisiae ADH1* gene is preferred.

The albumin is initially expressed with a secretion leader sequence, which may
30 be any leader effective in the yeast chosen. Leaders useful in *S. cerevisiae*

include that from the mating factor α polypeptide (MF α -1) and the hybrid leaders of EP-A-387 319. Such leaders (or signals) are cleaved by the yeast before the mature albumin is released into the surrounding medium. When the yeast strain lacks Kex2p activity (or equivalent) as well as being *yap3*, it may
5 be advantageous to choose a secretion leader which need not be cleaved from the albumin by Kex2p. Such leaders include those of *S. cerevisiae* invertase (*SUC2*) disclosed in JP 62-096086 (granted as 91/036516), acid phosphatase (*PHO5*), the pre-sequence of MF α -1, β -glucanase (*BGL2*) and killer toxin; *S. diastaticus* glucoamylase II; *S. carlsbergensis* α -galactosidase (*MEL1*); *K. lactis*
10 killer toxin; and *Candida* glucoamylase.

Various non-limiting embodiments of the invention will now be described by way of example and with reference to the accompanying drawings in which:

15 Figure 1 is a general scheme for the construction of mutated rHA expression plasmids, in which HA is a human albumin coding sequence, L is a sequence encoding a secretion leader, P is the PRB1 promoter, T is the ADH1 terminator, amp is an ampicillin resistance gene and LEU2 is the leucine selectable marker;

20

Figure 2 is a drawing representing a Western blot analysis of mutant rHA secreted by *S. cerevisiae*, in which Track A represents the culture supernatant from DB1 cir° pAYE316 (normal rHA), Track B represents the culture supernatant from DB1 cir+ pAYE464 (alteration 1), and Track C represents the
25 culture supernatant from DB1 cir+ pAYE468 (alteration 3);

Figure 3 is a scheme of the construction of pAYE515;

30 Figure 4 is a comparison of rHA fragment production by wild-type and protease-disrupted strains, presented as a drawing of an anti-HSA Western blot

of culture supernatant from shake flask cultures separated by non-reducing 10% SDS/PAGE, in which Track A corresponds to DB1 *cir°* pAYE316, Track B corresponds to DXY10 *cir°* pAYE316 (*yap3* strain), and Track C corresponds to ABB50 *cir°* pAYE316 (*yap3, kex2* strain);

5

Figure 5 is similar to Figure 4 but shows Coomassie Brilliant Blue stained 12.5% SDS Phastgel (Pharmacia) of culture supernatants from fed batch fermentations, namely Track D for the HSA standard, Track E for DB1 *cir°* pAYE316, Track F for DB1 *Δkex2 cir°* pAYE522, and Track G for DXY10 *cir°* pAYE522; and

10

Figure 6 is a scheme for the construction of pAYE519.

Detailed description of the invention

15

All standard recombinant DNA procedures are as described in Sambrook *et al* (1989) unless otherwise stated. The DNA sequences encoding HSA are derived from the cDNA disclosed in EP 201 239.

20 Example 1: Modification of the HSA cDNA.

In order to investigate the role of endoproteases in the generation of rHA fragment, the HSA cDNA (SEQ1 (which includes a sequence encoding the artificial secretion leader sequence of WO 90/01063)) was modified by site-directed mutagenesis. Three separate changes were made to the HSA sequence (SEQ2). The first, using the mutagenic primer FOG1, changed the Arg⁴¹⁰ codon only, replacing it with an Ala codon, leaving intact the dibasic site, Lys⁴¹³Lys⁴¹⁴. The second change, using primer FOG2, changed the residues 407-409, including the C-terminal residues of fragment, from LeuLeuVal to 25 AlaValAla. The third change, using the primer FOG3, altered residues 410-30

414 from ArgTyrThrLysLys (SEQ3) to AlaTyrThrGlnGln (SEQ4). The oligonucleotides encoded not only the amino acid changes, but also conservative base changes that create either a *Pvu*II or an *Spe*I restriction site in the mutants to facilitate detection of the changed sequences.

5

Single-stranded DNA of an M13mp19 clone, mp19.7 (EP 201 239; Figure 2), containing the HSA cDNA was used as the template for the mutagenesis reactions using the *In Vitro* Mutagenesis System, Version 2 (Amersham International plc) according to the manufacturer's instructions. Individual 10 plaques were selected and sequenced to confirm the presence of the mutations. Double stranded RF DNA was then made from clones with the expected changes and the DNA bearing the mutation was excised on an *Xba*I/*Sac*I fragment (Figure 1). This was used to replace the corresponding wild-type fragment of pAYE309 (EP 431 880; Figure 2). The presence of the mutated 15 *Xba*I/*Sac*I fragment within the plasmid was checked by digesting with *Pvu*II or *Spe*I as appropriate. These *Hind*III fragments were excised and inserted into the expression vector pAYE219 (Figure 1) to generate the plasmids pAYE464 (alteration 1, R410A), pAYE470 (alteration 2, L407A, L408V, V409A) and pAYE468 (alteration 3, R410A, K413Q, K414Q). These expression plasmids 20 comprise the *S. cerevisiae PRB1* promoter (WO 91/02057) driving expression of the HSA/MF α 1 leader sequence (WO 90/01063) fused in-frame with the mutated HA coding sequence which is followed by the *ADH1* transcription terminator. The plasmids also contain part of the 2 μ m plasmid to provide replication functions and the *LEU2* gene for selection of transformants.

25

pAYE464, pAYE470 and pAYE468 were introduced into *S. cerevisiae* DB1 cir⁺ (*a, leu2*; Sleep *et al*, 1990) by transformation and individual transformants were grown for 3 days at 30°C in 10ml YEPS (1% $^w/v$ yeast extract, 2% $^w/v$ peptone, 2% $^w/v$ sucrose) and then the supernatants were examined by anti-HSA 30 Western blot for the presence of the rHA fragment. The Western blots clearly

showed that fragment was still produced by the strains harbouring pAYE464, although the level was reduced slightly compared to the control expressing wild-type rHA. The mutations in the plasmid pAYE470 appeared to have no effect on the generation of fragment. However, DB1 cir⁺ pAYE468 showed
5 a novel pattern of HSA-related bands, with little or no fragment.

One example of each of DB1 cir⁺ pAYE464 and DB1 cir⁺ pAYE468 were grown to high cell density by fed batch culture in minimal medium in a fermenter (Collins, 1990). Briefly, a fermenter of 10L working volume was
10 filled to 5L with an initial batch medium containing 50 mL/L of a concentrated salts mixture (Table 1), 10 mL/L of a trace elements solution (Table 2), 50 mL/L of a vitamins mixture (Table 3) and 20 g/L sucrose. An equal volume of feed medium containing 100 mL/L of the salts mixture, 20 mL/L of the trace elements mixture, 100 mL/L of vitamins solution and 500 g/L sucrose
15 was held in a separate reservoir connected to the fermenter by a metering pump. The pH was maintained at 5.7 ± 0.2 by the automatic addition of ammonium hydroxide or sulphuric acid, and the temperature was maintained at 30°C. The stirrer speed was adjusted to give a dissolved oxygen tension of >20% air saturation at 1 v/v/min air flow rate.
20

Table 1. Salts Mixture

Chemical	Concentration (g/L)
KH ₂ PO ₄	114.0
MgSO ₄	12.0
CaCl ₂ .6H ₂ O	3.0
Na ₂ EDTA	2.0

Table 2. Trace Elements Solution

Chemical	Concentration (g/L)
ZnSO ₄ .7H ₂ O	3.0
FeSO ₄ .7H ₂ O	10.0
MnSO ₄ .4H ₂ O	3.2
CuSO ₄ .5H ₂ O	0.079
H ₃ BO ₃	1.5
KI	0.2
Na ₂ MoO ₄ .2H ₂ O	0.5
CoCl ₂ .6H ₂ O	0.56
H ₃ PO ₄	75mL/L

Table 3. Vitamins Solution

Chemical	Concentration (g/L)
Ca pantothenate	1.6
Nicotinic acid	1.2
m inositol	12.8
Thiamine HCl	0.32
Pyridoxine HCl	0.8
Biotin	0.008

The fermenter was inoculated with 100 mL of an overnight culture of *S. cerevisiae* grown in buffered minimal medium (Yeast nitrogen base [without amino acids, without ammonium sulphate, Difco] 1.7 g/L, (NH₄)₂SO₄ 5 g/L, citric acid monohydrate 6.09 g/L, Na₂HPO₄ 20.16 g/L, sucrose 20 g/L, pH6.5). The initial batch fermentation proceeded until the carbon source had been consumed, at which point the metering pump was switched on and the addition of feed was computer controlled (the micro MFCS system, B. Braun,

Melsungen, Germany) using an algorithm based on that developed by Wang *et al* (1979). A mass spectrometer was used in conjunction with the computer control system to monitor the off gases from the fermentation and to control the addition of feed to maintain a set growth rate (eg 0.1 h⁻¹). Maximum conversion of carbon substrate into biomass is achieved by maintaining the respiratory coefficient below 1.2 (Collins, 1990) and, by this means, cell densities of approximately 100 g/L cell dry weight can be achieved. The culture supernatants were compared with those of a wild-type rHA producer by Coomassie-stained SDS/PAGE and by Western blot. These indicated (Figure 2) that, whilst elimination of the monobasic Arg⁴¹⁰ (pAYE464) did reduce the level of the fragment by a useful amount, removal of both potential protease sites (pAYE468) almost abolished the 45kDa fragment.

The above data suggested that the generation of rHA fragment might be due to endoproteolytic attack, though the absence of an effect of removal of the potential Kex2p site Lys⁴¹³Lys⁴¹⁴ (Sleep *et al*, 1990, and confirmed by other studies not noted here) unless combined with elimination of Arg⁴¹⁰, had suggested a complex etiology. The reduction in the amount of fragment with the mutated rHA could in principle be due to an effect of the changes on the kinetics of folding of the molecule and not due to the removal of protease cleavage sites.

Example 2: Disruption of the *YAP3* gene.

The *YAP3* gene encoding yeast aspartyl protease 3 was mutated by the process of gene disruption (Rothstein 1983) which effectively deleted part of the *YAP3* coding sequence, thereby preventing the production of active Yap3p.

Four oligonucleotides suitable for PCR amplification of the 5' and 3' ends of the *YAP3* gene (Egel-Mitani *et al*, 1990) were synthesised using an Applied

Biosystems 380B Oligonucleotide Synthesiser. To assist the reader, we include as SEQ15 the sequence of the *YAP3* gene, of which 541-2250 is the coding sequence.

5' 5' end

YAP3A: 5' -CGTCAGACCTTGCATGCAGCCAAGACACCCTCACATAGC-3'

(SEQ5)

YAP3B: 5' -CCGTTACGTTCTGTGGTGGCATGCCACTTCCAAGTCCACCG-3'

(SEQ6)

10

3' end

YAP3C: 5'-GCGTCTCATAGTGGAAAAGCTTCTAAATACGACAACCTTCCCC-3'
(SEQ7)

YAP3D: 5'-CCCAAAATGGTACCTGTGTCATCACTCGTTGGGATAATACC-3'
(SEQ8)

15

PCR reactions were carried out to amplify individually the 5' and 3' ends of the *YAP3* gene from *S. cerevisiae* genomic DNA (Clontech Laboratories, Inc). Conditions were as follows: 2.5 µg/ml genomic DNA, 5 µg/ml of each primer, denature at 94°C 61 seconds, anneal at 37°C 121 secs, extend at 72°C 181 secs for 40 cycles, followed by a 4°C soak, using a Perkin-Elmer-Cetus Thermal Cycler and a Perkin-Elmer-Cetus PCR kit according to the manufacturer's recommendations. Products were analysed by gel electrophoresis and were found to be of the expected size. The 5' fragment was digested with *Sph*I and cloned into the *Sph*I site of pUC19HX (pUC19 lacking a *Hind*III site) to give pAYE511 (Figure 3), in which the orientation is such that *YAP3* would be transcribed towards the *Kpn*I site of the pUC19HX polylinker. The 3' *YAP3* fragment was digested with *Hind*III and *Asp*718 (an isoschizomer of *Kpn*I) and ligated into pUC19 digested with *Hind*III/*Asp*718 to give pAYE512. Plasmid DNA sequencing was carried out on the inserts to confirm that the desired sequences had been cloned. The *Hind*III/*Asp*718 fragment of pAYE512 was then subcloned into the corresponding sites of pAYE511 to give pAYE513 (Fig 3), in which the 5' and 3' regions of *YAP3*

are correctly orientated with a unique *Hind*III site between them. The *URA3* gene was isolated from YEp24 (Botstein *et al*, 1979) as a *Hind*III fragment and then inserted into this site to give pAYE515 (Fig 3), with *URA3* flanked by the 5' and 3' regions of *YAP3*, and transcribed in the opposite direction to *YAP3*.

5

A *ura3* derivative of strain DB1 *cir*^o pAYE316 (Sleep *et al*, 1991) was obtained by random chemical mutagenesis and selection for resistance to 5-fluoro-orotic acid (Boeke *et al*, 1987). The strain was grown overnight in 100 mL buffered minimal medium and the cells were collected by centrifugation and 10 then washed once with sterile water. The cells were then resuspended in 10 mL sterile water and 2 mL aliquots were placed in separate 15 mL Falcon tubes. A 5 mg/mL solution of N-methyl-N'-nitro-N-nitrosoguanidine (NTG) was then added to the tubes as follows: 0 μ L, 20 μ L, 40 μ L, 80 μ L or 160 μ L. The cells were then incubated at 30°C for 30 min and then centrifuged 15 and washed three times with sterile water. Finally, the cells were resuspended in 1 mL YEP (1% w/v yeast extract, 2% w/v Bacto peptone) and stored at 4°C. The percentage of cells that survived the mutagenic treatment was determined by spreading dilutions of the samples on YEP plates containing 2% w/v sucrose and incubating at 30°C for 3 days. Cells from the treatment which 20 gave approximately 50% survival were grown on YEP plates containing 2% w/v sucrose and then replica-plated onto YNB minimal medium containing 2% w/v sucrose and supplemented with 5-fluoro-orotic acid (1 mg/mL) and uracil (50 μ g/mL). Colonies able to grow on this medium were purified, tested to verify that they were unable to grow in the absence of uracil supplementation 25 and that this defect could be corrected by introduction of the *URA3* gene by transformation. One such strain, DBU3 *cir*^o pAYE316, was transformed with the *Sph*I/*Asp*718 *YAP3-URA3-YAP3* fragment of pAYE515 with selection for Ura⁺ colonies. A Southern blot of digested genomic DNA of a number of transformants was probed with the 5' and 3' ends of the *YAP3* gene and 30 confirmed the disruption of the *YAP3* gene. An anti-HSA Western blot of

YEPS shake-flask supernatants of two transformants indicated that disruption of *YAP3* markedly reduced rHA fragment levels.

One *yap3* derivative of DBU3 cir° pAYE316, designated DXY10 cir°
5 pAYE316, was grown several times by fed-batch fermentation in minimal
medium to high cell dry weight. When supernatants were examined by
Coomassie-stained PAGE and anti-HSA Western blot (Figs 4 and 5), the
reduction in the level of rHA 45kDa fragment was clearly apparent; estimates
of the amount of the degradation product vary from $\frac{1}{3}$ to $\frac{1}{5}$ of the levels seen
10 with the *YAP3* parent. The amount of rHA produced was not adversely
affected by the *yap3* mutation, indeed DXY10 cir° pAYE316 was found to
produce 30-50% more rHA than the *YAP3* equivalent, DB1 cir° pAYE316.
Despite the fact that cleavage of the leader sequence from the HA sequence is
C-terminal to a pair of basic residues, the rHA was found to have the correct
15 N-terminus.

The fermentation broth was centrifuged to remove the cells and then subject to
affinity chromatographic purification as follows. The culture supernatant was
passed through a Cibacron Blue F3GA Sepharose column (Pharmacia) which
20 was then washed with 0.1M phosphate glycine buffer, pH8.0. The rHA was
then eluted from the column with 2M NaCl, 0.1M phosphate glycine, pH8.0,
at which point it was >95% pure. It may be purified further by techniques
known in the art.

25 The albumin may alternatively be purified from the culture medium by any of
the variety of known techniques for purifying albumin from serum or
fermentation culture medium, for example those disclosed in WO 92/04367,
Maurel *et al* (1989), Curling (1980) and EP 524 681.

Example 3: Disruption of the *KEX2* gene in a *yap3* strain.

To construct a strain lacking both Yap3p and Kex2p activity, a *lys2* derivative of yeast strain DXY10 cir° (pAYE316) was obtained by random chemical mutagenesis and selection for resistance to α -amino adipate (Barnes and Thorner, 1985). Cells were mutagenised as in Example 2 and then plated on YNB minimal medium containing 2% w/v sucrose and supplemented with 2 mg/mL DL- α -amino adipate as the sole nitrogen source and 30 μ g/mL lysine. Colonies able to grow on this medium were purified and tested to verify that they were unable to grow in the absence of lysine supplementation and that this defect could be corrected by the introduction of the *LYS2* gene by transformation. This strain was then mutated by the process of gene disruption which effectively disrupted part of the *KEX2* coding sequence, thereby preventing production of active Kex2p. To assist the reader, the sequence of the *KEX2* gene is reproduced herein as SEQ14, of which 1329-3773 is the coding sequence.

Four oligonucleotides suitable for PCR amplification of the 5' and 3' ends of the *KEX2* gene (Fuller *et al*, 1989) were synthesised using an Applied Biosystems 380B Oligonucleotide Synthesiser.

5' end

KEX2A: 5'-CCATCTGGATCCAATGGTGCTTGGCCAAATAAATAGTTTCAGC-3'
25 KEX2B: 5'-GCTTCTTTACCGGTAACAAGCTTGAGTCCATTGG-3'
(SEQ9)
(SEQ10)

3' end

KEX2C: 5'-GGTAAGGTTAGTCGACCTATTTTGTTGTCTGC-3'
30 KEX2D: 5'-GGAAACGTATGAATTGATATCATTGATACAGACTCTGAGTACG-3'
(SEQ11)
(SEQ12)

PCR reactions were carried out to amplify individually the 5' and 3' ends of the *KEX2* gene from *S. cerevisiae* genomic DNA (Clontech Laboratories Inc). Conditions were as follows: 2.5 µg/ml genomic DNA, 5 µg/ml of each primer, denature 94°C 61s, anneal 37°C 121s, extend 72°C 181s for 40 cycles, followed by a 4°C soak, using a Perkin-Elmer-Cetus Thermal Cycler and a Perkin-Elmer-Cetus PCR kit according to the manufacturer's recommendations. Products were analysed by gel electrophoresis and were found to be of the expected size (0.9 kb for the 5' product and 0.62 kb for the 3' product). The 5' product was digested with *Bam*HI and *Hind*III and the 3' product was digested with *Hind*III and *Sal*I and then the two fragments were together cloned into pUC19HX digested with *Bam*HI and *Sal*I. A 4.8 kb *Hind*III fragment comprising the *S. cerevisiae LYS2* gene (Barnes & Thorner, 1985) was then inserted into the resulting plasmid at *Hind*III (ie between the two *KEX2* fragments) to form pAYE519 (Fig 6).

15

The *lys2* derivative of DXY10 cir° (pAYE316), *lys2-16*, was transformed with the 6.0 kb *KEX2-LYS2-KEX2* fragment of pAYE519, selecting for Lys⁺ colonies. A Southern blot of digested genomic DNA of a number of transformants was probed with the 5' and 3' ends of the *KEX2* gene and confirmed the disruption of the *KEX2* gene. An anti-HSA Western blot of YEPS shake-flask culture supernatants of these transformants indicated that disruption of *KEX2* in a *yap3* strain reduced the level of rHA fragment still further, despite the lack of an effect of disruption of *KEX2* alone in Example 4 below. Analysis of the rHA produced by one such strain, ABB50, indicated that the leader sequence was incorrectly processed, leading to an abnormal N-terminus.

30 The strain ABB50 (pAYE316) was cured of its plasmid (Sleep *et al*, 1991) and transformed with a similar plasmid, pAYE522, in which the hybrid leader sequence was replaced by the *S. cerevisiae* invertase (*SUC2*) leader sequence

such that the encoded leader and the junction with the HSA sequence were as follows:

5 MLLQ AFLF LLAGFAAKISA ↓ DAHKS (SEQ13)
Invertase leader HSA

In this construct, cleavage of the leader sequence from HSA does not rely upon activity of the Kex2 protease. The strain ABB50 (pAYE522) was found to produce rHA with a similarly very low level of rHA fragment, but in this instance the N-terminus corresponded to that of serum-derived HSA, ie there was efficient and precise removal of the leader sequence.

Example 4: Disruption of the *KEX2* gene alone (Comparative Example).

15 By a similar method to that disclosed in Example 3 the *KEX2* gene was disrupted in *S. cerevisiae*. This strain had the Yap3p proteolytic activity and was therefore not within the scope of the invention. When this strain was grown in fed batch fermentation the rHA produced contained similar amounts of fragment to that produced by strains with an intact *KEX2* gene. In addition,
20 the overall level of rHA was reduced and the leader sequence was not correctly processed, leading to an abnormal N-terminus.

Example 5: Identification of equivalent protease in *Pichia*.

25 As noted above, non-*Saccharomyces* yeast similarly produce the undesirable
fragment of rHA and therefore have the Yap3p proteolytic activity. We have
confirmed this by performing Southern hybridisations of *Pichia angusta* DNA,
using the *S. cerevisiae* *YAP3* gene as a probe. A specific DNA fragment was
identified, showing that, not only is the Yap3p proteolytic activity present in
30 *P. angusta*, but a specific homologue of the *YAP3* gene is present also.

The method of Southern hybridization used for detection of the *YAP3* homologue can be adapted to clone the gene sequence from a genomic DNA library of *Pichia* DNA using standard procedures (Sambrook *et al*, 1989). Disruption of the *YAP3* homologue in *Pichia sp.* can be achieved using similar
5 techniques to those used above for *Saccharomyces* (Cregg and Madden, 1987).

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SEQUENCE LISTING

(1) GENERAL INFORMATION:

(i) APPLICANT:

- (A) NAME: Delta Biotechnology Limited
- (B) STREET: Castle Court, Castle Boulevard
- (C) CITY: Nottingham
- (D) STATE: Nottinghamshire
- (E) COUNTRY: United Kingdom
- (F) POSTAL CODE (ZIP): NG7 1FD

(ii) TITLE OF INVENTION: Yeast strains and modified albumins

(iii) NUMBER OF SEQUENCES: 15

(iv) COMPUTER READABLE FORM:

- (A) MEDIUM TYPE: Floppy disk
- (B) COMPUTER: IBM PC compatible
- (C) OPERATING SYSTEM: PC-DOS/MS-DOS
- (D) SOFTWARE: PatentIn Release #1.0, Version #1.25 (EPO)

(2) INFORMATION FOR SEQ ID NO: 1:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1830 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA to mRNA

(iii) HYPOTHETICAL: NO

(iv) ANTI-SENSE: NO

(vi) ORIGINAL SOURCE:

- (A) ORGANISM: Homo sapiens

(ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) LOCATION: 73..1827

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 1:

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Asp Leu Gly Glu Asn Phe Lys Ala Leu Val Leu Ile Ala Phe Ala	
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CAG TAT CTT CAG CAG TGT CCA TTT GAA GAT CAT GTA AAA TTA GTG AAT	204
Gln Tyr Leu Gln Gln Cys Pro Phe Glu Asp His Val Lys Leu Val Asn	
30 35 40	
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45 50 55 60	

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320	325	330	
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(2) INFORMATION FOR SEQ ID NO: 2:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 585 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 2:

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Glu	Asn	Phe	Lys	Ala	Leu	Val	Leu	Ile	Ala	Phe	Ala	Gln	Tyr	Leu	Gln
		20						25					30		
Gln	Cys	Pro	Phe	Glu	Asp	His	Val	Lys	Leu	Val	Asn	Glu	Val	Thr	Glu
	35					40					45				
Phe	Ala	Lys	Thr	Cys	Val	Ala	Asp	Glu	Ser	Ala	Glu	Asn	Cys	Asp	Lys
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Ser	Leu	His	Thr	Leu	Phe	Gly	Asp	Lys	Leu	Cys	Thr	Val	Ala	Thr	Leu
	65				70				75			80			
Arg	Glu	Thr	Tyr	Gly	Glu	Met	Ala	Asp	Cys	Cys	Ala	Lys	Gln	Glu	Pro
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Glu	Arg	Asn	Glu	Cys	Phe	Leu	Gln	His	Lys	Asp	Asp	Asn	Pro	Asn	Leu
	100					105				110					
Pro	Arg	Leu	Val	Arg	Pro	Glu	Val	Asp	Val	Met	Cys	Thr	Ala	Phe	His
	115					120				125					
Asp	Asn	Glu	Glu	Thr	Phe	Leu	Lys	Lys	Tyr	Leu	Tyr	Glu	Ile	Ala	Arg
	130					135				140					
Arg	His	Pro	Tyr	Phe	Tyr	Ala	Pro	Glu	Leu	Leu	Phe	Phe	Ala	Lys	Arg
	145				150				155			160			
Tyr	Lys	Ala	Ala	Phe	Thr	Glu	Cys	Cys	Gln	Ala	Ala	Asp	Lys	Ala	Ala
	165					170				175					
Cys	Leu	Leu	Pro	Lys	Leu	Asp	Glu	Leu	Arg	Asp	Glu	Gly	Lys	Ala	Ser
	180					185				190					
Ser	Ala	Lys	Gln	Arg	Leu	Lys	Cys	Ala	Ser	Leu	Gln	Lys	Phe	Gly	Glu
	195					200				205					
Arg	Ala	Phe	Lys	Ala	Trp	Ala	Val	Ala	Arg	Leu	Ser	Gln	Arg	Phe	Pro
	210					215				220					
Lys	Ala	Glu	Phe	Ala	Glu	Val	Ser	Lys	Leu	Val	Thr	Asp	Leu	Thr	Lys
	225				230				235			240			
Val	His	Thr	Glu	Cys	Cys	His	Gly	Asp	Leu	Leu	Glu	Cys	Ala	Asp	Asp
	245					250				255					
Arg	Ala	Asp	Leu	Ala	Lys	Tyr	Ile	Cys	Glu	Asn	Gln	Asp	Ser	Ile	Ser
	260					265				270					
Ser	Lys	Leu	Lys	Glu	Cys	Cys	Glu	Lys	Pro	Leu	Leu	Glu	Lys	Ser	His
	275					280				285					
Cys	Ile	Ala	Glu	Val	Glu	Asn	Asp	Glu	Met	Pro	Ala	Asp	Leu	Pro	Ser

290	295	300
Leu Ala Ala Asp Phe Val Glu Ser Lys Asp Val Cys Lys Asn Tyr Ala		
305	310	315
320		
Glu Ala Lys Asp Val Phe Leu Gly Met Phe Leu Tyr Glu Tyr Ala Arg		
325	330	335
Arg His Pro Asp Tyr Ser Val Val Leu Leu Leu Arg Leu Ala Lys Thr		
340	345	350
Tyr Glu Thr Thr Leu Glu Lys Cys Cys Ala Ala Asp Pro His Glu		
355	360	365
Cys Tyr Ala Lys Val Phe Asp Glu Phe Lys Pro Leu Val Glu Glu Pro		
370	375	380
Gln Asn Leu Ile Lys Gln Asn Cys Glu Leu Phe Glu Gln Leu Gly Glu		
385	390	395
400		
Tyr Lys Phe Gln Asn Ala Leu Leu Val Arg Tyr Thr Lys Lys Val Pro		
405	410	415
Gln Val Ser Thr Pro Thr Leu Val Glu Val Ser Arg Asn Leu Gly Lys		
420	425	430
Val Gly Ser Lys Cys Cys Lys His Pro Glu Ala Lys Arg Met Pro Cys		
435	440	445
Ala Glu Asp Tyr Leu Ser Val Val Leu Asn Gln Leu Cys Val Leu His		
450	455	460
Glu Lys Thr Pro Val Ser Asp Arg Val Thr Lys Cys Cys Thr Glu Ser		
465	470	475
480		
Leu Val Asn Arg Arg Pro Cys Phe Ser Ala Leu Glu Val Asp Glu Thr		
485	490	495
Tyr Val Pro Lys Glu Phe Asn Ala Glu Thr Phe Thr Phe His Ala Asp		
500	505	510
Ile Cys Thr Leu Ser Glu Lys Glu Arg Gln Ile Lys Lys Gln Thr Ala		
515	520	525
Leu Val Glu Leu Val Lys His Lys Pro Lys Ala Thr Lys Glu Gln Leu		
530	535	540
Lys Ala Val Met Asp Asp Phe Ala Ala Phe Val Glu Lys Cys Cys Lys		
545	550	555
560		
Ala Asp Asp Lys Glu Thr Cys Phe Ala Glu Glu Gly Lys Lys Leu Val		
565	570	575
Ala Ala Ser Gln Ala Ala Leu Gly Leu		
580	585	

(2) INFORMATION FOR SEQ ID NO: 3:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 5 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: peptide

(iii) HYPOTHETICAL: NO
(iii) ANTI-SENSE: NO
(v) FRAGMENT TYPE: internal

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 3:

Arg Tyr Thr Lys Lys
1 5

(2) INFORMATION FOR SEQ ID NO: 4:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 5 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(iii) HYPOTHETICAL: NO

(iii) ANTI-SENSE: NO

(v) FRAGMENT TYPE: internal

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 4:

Ala Tyr Thr Gln Gln
1 5

(2) INFORMATION FOR SEQ ID NO: 5:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 39 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iii) HYPOTHETICAL: NO

(iii) ANTI-SENSE: NO

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 5:

CGTCAGACCT TGCATGCAGC CAAGACACCC TCACATAGC

39

(2) INFORMATION FOR SEQ ID NO: 6:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 42 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iii) HYPOTHETICAL: NO
(iii) ANTI-SENSE: YES

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 6:

CCGTTACGTT CTGTGGTGGC ATGCCCACTT CCAAGTCCAC CG

42

(2) INFORMATION FOR SEQ ID NO: 7:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 42 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iii) HYPOTHETICAL: NO

(iii) ANTI-SENSE: NO

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 7:

GCGTCTCATA GTGGAAAAGC TTCTAAATAC GACAACTTCC CC

42

(2) INFORMATION FOR SEQ ID NO: 8:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 41 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iii) HYPOTHETICAL: NO

(iii) ANTI-SENSE: YES

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 8:

CCCAAAATGG TACCTGTGTC ATCACTCGTT GGGATAATAC C

41

(2) INFORMATION FOR SEQ ID NO: 9:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 44 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iii) HYPOTHETICAL: NO

(iii) ANTI-SENSE: NO

33

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 9:

CCATCTGGAT CCAATGGTGC TTTGCCAAA TAAATAGTTT CAGC

44

(2) INFORMATION FOR SEQ ID NO: 10:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 35 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iii) HYPOTHETICAL: NO

(iii) ANTI-SENSE: YES

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 10:

GCTTCCTTTA CCGGTAACAA GCTTGAGTCC ATTGG

35

(2) INFORMATION FOR SEQ ID NO: 11:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 37 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iii) HYPOTHETICAL: NO

(iii) ANTI-SENSE: NO

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 11:

GGTAAGGTTT AGTCGACCTA TTTTTGTTT TGTCTGC

37

(2) INFORMATION FOR SEQ ID NO: 12:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 44 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iii) HYPOTHETICAL: NO

(iii) ANTI-SENSE: YES

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 12:

GGAAACGTAT GAATTCGATA TCATTGATAC AGACTCTGAG TACG

44

(2) INFORMATION FOR SEQ ID NO: 13:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 24 amino acids
 (B) TYPE: amino acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(iii) HYPOTHETICAL: NO

(iii) ANTI-SENSE: NO

(v) FRAGMENT TYPE: N-terminal

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 13:

Met	Leu	Leu	Gln	Ala	Phe	Leu	Phe	Leu	Leu	Ala	Gly	Phe	Ala	Ala	Lys
1					5				10				15		
Ile	Ser	Ala	Asp	Ala	His	Lys	Ser								
					20										

(2) INFORMATION FOR SEQ ID NO: 14:

(i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 4106 base pairs
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iii) HYPOTHETICAL: NO

(iii) ANTI-SENSE: NO

(vi) ORIGINAL SOURCE:
 (A) ORGANISM: *Saccharomyces cerevisiae*

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 14:

GAATTCTCTG	TTGACTACTA	AACTGAGAGA	ATTTGCCGAG	ACTCTAAGAA	CAGCTTGAA	60
AGAGCGTTCT	GCCGATGATT	CCATAATTGT	CACTCTGAGA	GAGCAAATGC	AAAGAGAAAT	120
CTTCAGGTTG	ATGTCGTTGT	TCATGGACAT	ACCTCCAGTG	CAACCAAACG	AGCAATTCAC	180
TTGGGAATAC	GTTGACAAAG	ACAAGAAAAT	CCACACTATC	AAATCGACTC	CGTTAGAATT	240
TGCCTCCAAA	TACGCAAAAT	TGGACCCCTTC	CACGCCAGTC	TCATTGATCA	ATGATCCAAG	300
ACACCATATG	GTAAATTAAT	TAAGATCGAT	CGTTTAGGAA	ACGTCCCTTGG	CGGAGATGCC	360
GTGATTTACT	TAAATGTTGA	CAATGAAACA	CTATCTAAAT	TGGTTGTTAA	GAGATTACAA	420
AATAACAAAG	CTGTCTTTTT	TGGATCTCAC	ACTCCAAAGT	TCATGGACAA	GAAAACGTGGT	480
GTCATGGATA	TTGAATTGTG	GAACTATCCT	GCCATGGCTA	TAATTTACCT	CAGCAAAAGG	540
CATCCGGTAT	TAGATACCAT	GAAAGTTGA	TGACTCATGC	TATGTTGGAT	CACTGGCTGC	600
CACGTCGATG	AAACGTCTAA	ATTACCACTT	CGCTACCGTC	TGAAAATTCC	TGGGGTAAAG	660

ACTCCGGTAA AGACGGATT A TACGTGATGA CTCAAAAGTA CTTCGAGGAG TACTGCTTTC	720
AAATTGTGGT CGATATCAAT GAATTGCCAA AAGAGCTGGC TTCAAAATTC ACCTCAGGTA	780
AGGAAGAGCC GATTGTCTT G CCCATCTGG A CCCAATGGTG CTTGGCCAA ATAAATAGTT	840
TCAGCAGCTC TGATGTAGAT ACACGTATCT CGACATGTT TATTTTACT ATACATACAT	900
AAAAGAAATA AAAATGATA ACGTGTATAT TATTATTCAT ATAATCAATG AGGGTCATT	960
TCTGAAACGC AAAAACGGT AAATGGAAAA AAAATAAAGA TAGAAAAAGA AAACAAACAA	1020
AGGAAAGGTT AGCATATTAA ATAATGAGC TGATACTTC A ACAGCATCGC TGAAGAGAAC	1080
AGTATTGAAA CCGAACATT TTCTAAAGGC AAACAAGGT A CTCCATATTT GCTGGACGTG	1140
TTCTTCTCT CGTTTCATAT GCATAATTCT GTCATAAGCC TGTTCTTTT CCTGGCTTAA	1200
ACATCCC GTT TTGTAAGA GAAATCTATT CCACATATT CATTCAATTG GCTACCATAC	1260
TAAGGATAAA CTAATCCC GTT TGTTTTGG CCTCGTCACA TAATTATAAA CTACTAACCC	1320
ATTATCAGAT GAAAGTGAGG AAATATATTA CTTTATGCTT TTGGTGGGCC TTTCAACAT	1380
CCGCTCTGT ATCATCACAA CAAATCCAT TGAAGGACCA TACGTACGA CAGTATTTG	1440
CTGTAGAAAG CAATGAAACA TTATCCC GTT TGGAGGAAAT GCATCCAAT TGGAAATATG	1500
AACATGATGT TCGAGGGCTA CCAAACCAATT ATGTTTTTC AAAAGAGTTG CTAAAATTGG	1560
GCAAAAGATC ATCATTAGAA GAGTTACAGG GGGATAACAA CGACCACATA TTATCTGTCC	1620
ATGATTATT CCCCGTAAAC GACCTATT A AGAGACTACC GGTGCCTGCT CCACCAATGG	1680
ACTCAAGCTT GTTACCGGT A AAGAAGCTG AGGATAAACT CAGCATAAT GATCCGCTT	1740
TTGAGAGGCA GTGGCACTTG GTCAATCCAA GTTTCTGG CAGTGTATA AATGTTCTG	1800
ATCTGTGGTA CAATAATATT ACAGGCGCAG GGGTCGTGGC TGCCATTGTT GATGATGGCC	1860
TTGACTACGA AAATGAAGAC TTGAAGGATA ATTTTGCGC TGAAGGTTCT TGGGATTCA	1920
ACGACAATAC CAATTTACCT AAACCAAGAT TATCTGATGA CTACCATGGT ACGAGATGTG	1980
CAGGTGAAAT AGCTGCCAA AAAGGTAAACA ATTTTGCGG TGTGGGGTA GGTTACAACG	2040
CTAAAATCTC AGGCATAAGA ATCTTATCCG GTGATATCAC TACGGAAAGAT GAAGCTGCGT	2100
CCTTGATT A TGGTCTAGAC GTAAACGATA TATATTCA TCTCATGGGT CCCGCTGATG	2160
ACGGAAAGACA TTTACAAGGC CCTAGTGACC TGGTAAAAA GGCTTAGTA AAAGGTGTTA	2220
CTGAGGGAAAG AGATTCCAA GGAGCGATT ACGTTTTGC CAGTGGAAAT GGTGGAAC	2280
GTGGTGATAA TTGCAATTAC GACGGCTATA CTAATTCCAT ATATTCTATT ACTATTGGG	2340
CTATTGATCA CAAAGATCTA CATCCTCCTT ATTCCGAAGG TTGTTCCGCC GTCATGGCAG	2400
TCACGTATTC TTCAGGTTCA GGCAGATATA TTCATTGAG TGATATCAAC GGCAGATGCA	2460
GTAATAGCCA CGGTGGAACG TCTGCGGCTG CTCCATTAGC TGCCGGTGT TACACTTTGT	2520
TACTAGAAGC CAACCCAAAC CTAACCTGGA GAGACGTACA GTATTTATCA ATCTGTCTG	2580
CGGTAGGGTT AGAAAAGAAC GCTGACGGAG ATTGGAGAGA TAGGCCATG GGGAAAGAAAT	2640

ACTCTCATCG CTATGGCTTT GGTAAAATCG ATGCCATAA GTTAATTGAA ATGTCCAAGA	2700
CCTGGGAGAA TGTAAACGCA CAAACCTGGT TTTACCTGCC AACATTGTAT GTTCCCAGT	2760
CCACAAACTC CACGGAAAGAG ACATTAGAAT CCGTCATAAC CATATCAGAA AAAAGTCTTC	2820
AAGATGCTAA CTTCAAGAGA ATTGAGCACG TCACGGTAAC TGTAGATATT GATACAGAAA	2880
TTAGGGGAAC TACGACTGTC GATTTAATAT CACCAGCGGG GATAATTCA AACCTTGGCG	2940
TTGTAAGACC AAGAGATGTT TCATCAGAGG GATTCAAAGA CTGGACATTG ATGTCTGTAG	3000
CACATTGGGG TGAGAACGGC GTAGGTGATT GGAAAATCAA GTTAAAGACA ACAGAAAATG	3060
GACACAGGAT TGACTTCCAC AGTTGGAGGC TGAAGCTCTT TGGGAATCC ATTGATTCA	3120
CTAAAACAGA AACCTTCGTC TTTGGAAACG ATAAAGAGGA GTTGAACCA GCTGCTACAG	3180
AAAGTACCGT ATCACAATAT TCTGCCAGTT CAACTTCTAT TTCCATCAGC GCTACTTCTA	3240
CATCTTCTAT CTCAATTGGT GTGGAAACGT CGGCCATTCC CCAAACGACT ACTGCGAGTA	3300
CCGATCCTGA TTCTGATCCA AACACTCCTA AAAAACTTTC CTCTCCTAGG CAAGCCATGC	3360
ATTATTTTT ACAAATATTT TTGATTGGCG CCACATTTT GGTGTTATAC TTCATGTTTT	3420
TTATGAAATC AAGGAGAAGG ATCAGAAGGT CAAGAGCGGA AACGTATGAA TTCGATATCA	3480
TTGATACAGA CTCTGAGTAC GATTCTACTT TGGACAATGG AACTTCCGGA ATTACTGAGC	3540
CCGAAGAGGT TGAGGACTTC GATTTGATT TGTCCGATGA AGACCATCTT GCAAGTTGT	3600
CTTCATCAGA AAACGGTGAT GCTGAACATA CAATTGATAG TGTACTAACAAACGAAAATC	3660
CATTTAGTGA CCCTATAAAG CAAAAGTTCC CAAATGACGC CAACGCAGAA TCTGCTTCCA	3720
ATAAAATTACA AGAATTACAG CCTGATGTTG CTCCATCTTC CGGACGATCG TGATTCGATA	3780
TGTACAGAAA GCTTCAAATT ACAAAATAGC ATTTTTTCT TATAGATTAT AATACTCTCT	3840
CATACGTATA CGTATATGTG TATATGATAT ATAAACAAAC ATTAATATCC TATTCTTCC	3900
GTTTGAAATC CCTATGATGT ACTTTGCATT GTTGCACCC GCGAATAAAA TGAAAACCTCC	3960
GAACCGATAT ATCAAGCACA TAAAAGGGGA CGGTCCAATT AATGCATATT TAAGACCACA	4020
GCTGAATAAC TTTAAAACGG CAGACAAAAC AAAAATAGG TCGAATAAAC CTTACCTGCC	4080
TAGAAGGAAT GACAGCAGCT AATAAG	4106

(2) INFORMATION FOR SEQ ID NO: 15:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 2526 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iii) HYPOTHETICAL: NO

(iii) ANTI-SENSE: NO

- (vi) ORIGINAL SOURCE:
 - (A) ORGANISM: *Saccharomyces cerevisiae*

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 15:

CCGTTTTCTT	TTCGTAAAAA	AAAACAATAG	ACACTATATA	TAGACACTTT	TTCCTTTCTT	60
TCTTGCGCG	ATTCAAGAG	GAAAAGCATA	CTTAAATAAG	AATATTCTA	AAACACACGT	120
TCTGACGCGT	CAATTAGATC	GTCAGACCTT	GCATGCAGCC	AAGACACCCCT	CACATAGCAC	180
TGCCTCCTTC	CTCCTCTTT	CTGTCACCAC	CTCACCTCCC	TCGTCCACTC	AACTGAGTGG	240
CTTTTCGCTC	CTTTTATACT	GCGCCATGAG	TAGTTTCGT	TTCACTGATG	TGTCCGAAAA	300
AATTGAGGTT	TCATAAAAAA	ATTCGTGGAC	TTATTTATGG	AGAAACAGGG	AAATCCGACT	360
ACTTAAGAAA	AGGGTGTCAA	AGAGGATTAA	CTTTTTCTT	TCTTTTGCA	TTTGTTCCTA	420
TTTCCGCAAT	TGGACGGTTA	TTAAGAAGAA	CGCAATTGGC	TTTTCTGTAT	ATTAAAATAC	480
ATAGCGTAAT	AAAAAGATAA	GGTGAACACC	AAGCATATAG	TATAATATTA	CCTACCACAT	540
ATGAAACTGA	AAACTGTAAG	ATCTGCGTC	CTTCGTCAC	TCTTGCATC	GCAGGTTCTC	600
GGTAAGATAA	TACCAGCAGC	AAACAAGCGC	GACGACGACT	CGAATTCCAA	GTTCGTCAAG	660
TTGCCCTTTC	ATAAGCTTTA	CGGGGACTCG	CTAGAAAATG	TGGGAAGCGA	CAAAAAACCG	720
GAAGTACGCC	TATTGAAGAG	GGCTGACGGT	TATGAAGAAA	TTATAATTAC	CAACCAGCAA	780
AGTTTCTATT	CGGTGGACTT	GGAAAGTGGGC	ACGCCACCAC	AGAACGTAAC	GGTCCTGGTG	840
GACACAGGCT	CCTCTGATCT	ATGGATTATG	GGCTCGGATA	ATCCATACTG	TTCTTCGAAAC	900
AGTATGGGTA	GTAGCCGGAG	ACGTGTTATT	GACAAACGTG	ATGATTGTC	AAGCGGCCGA	960
TCTTGATTAA	ATGATATAAA	CCCATTTGGC	TGGTTGACGG	GAACGGGCAG	TGCCATTGGC	1020
CCCACTGCTA	CGGGCTTAGG	AGGCGGTTCA	GGTACGGCAA	CTCAATCCGT	GCCTGCTTCG	1080
GAAGCCACCA	TGGACTGTCA	ACAATACGGG	ACATTTCCA	CTTCGGGCTC	TTCTACATT	1140
AGATCAAACA	ACACCTATTT	CAGTATTAGC	TACGGTGATG	GGACTTTGC	CTCCGGTACT	1200
TTTGGTACGG	ATGTTTGGA	TTTAAGCGAC	TTGAACGTTA	CCGGGTTGTC	TTTGCCGTT	1260
GCCAATGAAA	CGAATTCTAC	TATGGGTGTG	TTAGGTATTG	GTGCCCCGA	ATTAGAAGTC	1320
ACTTATTCTG	GCTCTACTGC	GTCTCATAGT	GGAAAAGCTT	ATAAAATACGA	CAACTTCCCC	1380
ATTGTATTGA	AAAATTCTGG	TGCTATCAAA	AGCAACACAT	ATTCTTGTA	TTGAACGAC	1440
TCGGACGCTA	TGCATGGCAC	CATTTGTTC	GGAGCCGTGG	ACCACAGTAA	ATATACCGGC	1500
ACCTTATACA	CAATCCCCAT	CGTAAACACT	CTGAGTGCTA	GTGGATTAG	CTCTCCCATT	1560
CAATTGATG	TCACTATTAA	TGGTATCGGT	ATTAGTGATT	CTGGGAGTAG	TAACAAGACC	1620
TTGACTACCA	CTAAAATACC	TGCTTTGTG	GATTCCGGTA	CTACTTTGAC	TTATTTACCT	1680
CAAACAGTGG	TAAGTATGAT	CGCTACTGAA	CTAGGGTGCAC	AATACTCTTC	CAGGATAGGG	1740
TATTACGTAT	TGGACTGTCC	ATCTGATGAT	AGTATGGAAA	TAGTGTTCGA	TTTGCGTGGT	1800
TTTCACATCA	ATGCACCACT	TTCGAGTTT	ATCTTGAGTA	CTGGCACTAC	ATGTCTTTA	1860
GGTATTATCC	CAACGAGTGA	TGACACAGGT	ACCATTTGG	GTGATTCAATT	TTTGACTAAC	1920

GCGTACGTGG TTTATGATTT GGAGAATCTT GAAATATCCA TGGCACAAGC TCGCTATAAT	1980
ACCACAAGCG AAAATATCGA AATTATCACA TCCTCTGTTCAAGCGCCGT AAAGGCACCA	2040
GGCTATACAA ACACCTGGTC CACAAGTGCA TCTATTGTTA CCGGTGGTAA CATATTTACT	2100
GTAAATTCTC CACAAACTGC TTCCCTTAGC GGTAACCTGA CGACCAGTAC TGCACTCCGCC	2160
ACTTCTACAT CAAGTAAAAG AAATGTTGGT GATCATATAG TTCCATCTTT ACCCCTCACA	2220
TTAATTCTC TTCTTTTGC ATTCACTGAA AACCGTTGC ACAAAAGTTA GACATTCA	2280
TCTCCAAGCC AGTTGGAGTT TCTGGCGGAA ATCGTTGCTC TCGCTGGC AAAGTTTTT	2340
TTTATTATTA ATTTTTTATT GTTACGTTGG CGGTCTTTAT TTTTACTTCA CAATAGTTA	2400
TCTTACCCAC TAAGAATAGG TTACCATTAA TTCACATTT TTTTCTCAT TCCTAGTATA	2460
CTATTTACCT GGGATATGGC CTATAATCAA AGGCTTAAT ATTCTAATAA TTCGTTGGC	2520
ATCTAG	2526

CLAIMS

1. A process for preparing albumin by secretion from a yeast genetically modified to produce and secrete the albumin, comprising culturing the yeast in a culture medium such that albumin is secreted into the medium, characterised in that the yeast cells have a reduced level of yeast aspartyl protease 3 proteolytic activity.
5
2. A process according to Claim 1 wherein the said proteolytic activity is an endoprotease activity specific for monobasic sites and for paired basic amino acids in a polypeptide.
10
3. A process according to Claim 1 or 2 wherein the yeast is *S. cerevisiae*.
15
4. A process according to Claim 1, 2 or 3 wherein the yeast lacks a functional *YAP3* gene or homologue thereof.
5. A process according to any one of Claims 1 to 4 wherein the yeast cells additionally have a reduced level of *S. cerevisiae* Kex2p proteolytic activity.
20
6. A process according to any one of the preceding claims wherein the albumin is a human albumin.
25
7. A culture of yeast cells containing a polynucleotide sequence encoding an albumin and a second polynucleotide sequence encoding a secretion signal causing albumin expressed from the first polynucleotide sequence to be secreted from the yeast, characterised in that the yeast cells have a reduced level of yeast aspartyl protease 3 proteolytic
30

activity.

8. A culture according to Claim 7 wherein the albumin is a human albumin.

5

9. A culture according to Claim 7 or 8 wherein the yeast is *S. cerevisiae*.

10. A culture according to any one of Claims 7 to 9 wherein the said signal is cleaved by the yeast prior to release of the albumin from the yeast.

11. A culture according to any one of Claims 7 to 10 wherein the yeast cells additionally have a reduced level of Kex2p proteolytic activity.

15

12. A culture according to Claim 11 wherein the said secretion signal is cleaved from the albumin by a protease other than Kex2p.

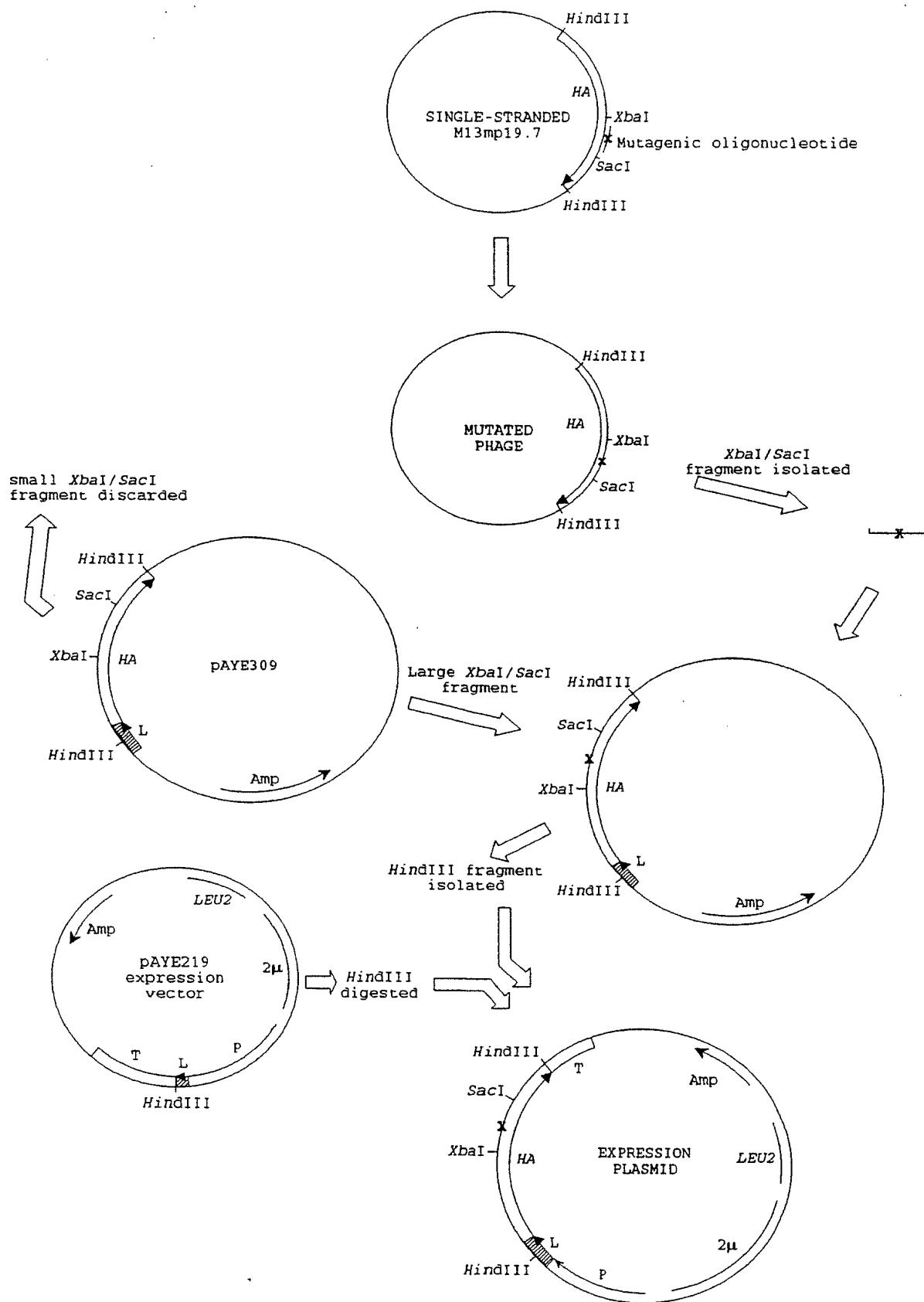
13. A modified albumin having at least 90% sequence identity to a naturally-occurring albumin, which naturally-occurring albumin is susceptible to cleavage with yeast aspartyl protease 3 (Yap3p) when expressed and secreted in yeast, characterised in that the modified albumin is not susceptible to such cleavage.

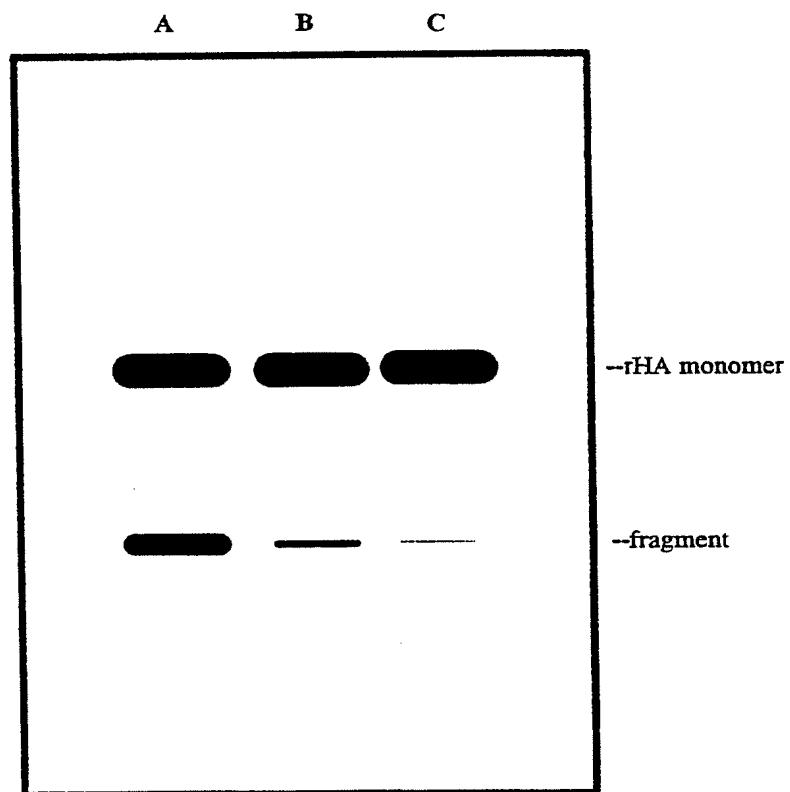
- 25 14. A modified albumin according to Claim 13 wherein the modified albumin lacks a monobasic amino acid present in the naturally-occurring albumin protein.

- 30 15. A modified albumin according to Claim 13 or 14 wherein the said monobasic amino acid is arginine.

16. A modified albumin according to Claim 14 or 15 wherein the modified albumin additionally lacks a pair of basic amino acids present in the naturally-occurring albumin.
- 5 17. A modified albumin according to Claim 16 wherein the said pair of amino acids is Lys, Lys; Lys, Arg; Arg, Lys; or Arg, Arg.
- 10 18. A modified albumin according to Claim 13 wherein the naturally- occurring albumin is a human albumin and the modified protein lacks Arg⁴¹⁰; and, optionally, residues 413 and 414 do not each consist of lysine or arginine.
- 15 19. A modified albumin according to Claim 18 which is a human albumin having the amino acid changes R410A, K413Q, K414Q.
- 20 20. A polynucleotide encoding a modified albumin according to any one of Claims 13 to 19.
21. A yeast containing a polynucleotide according to Claim 20, transcription signals such that the modified albumin is expressed in the yeast, and a further polynucleotide adjacent the said polynucleotide such that the modified albumin is secreted from the yeast.

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Figure 1

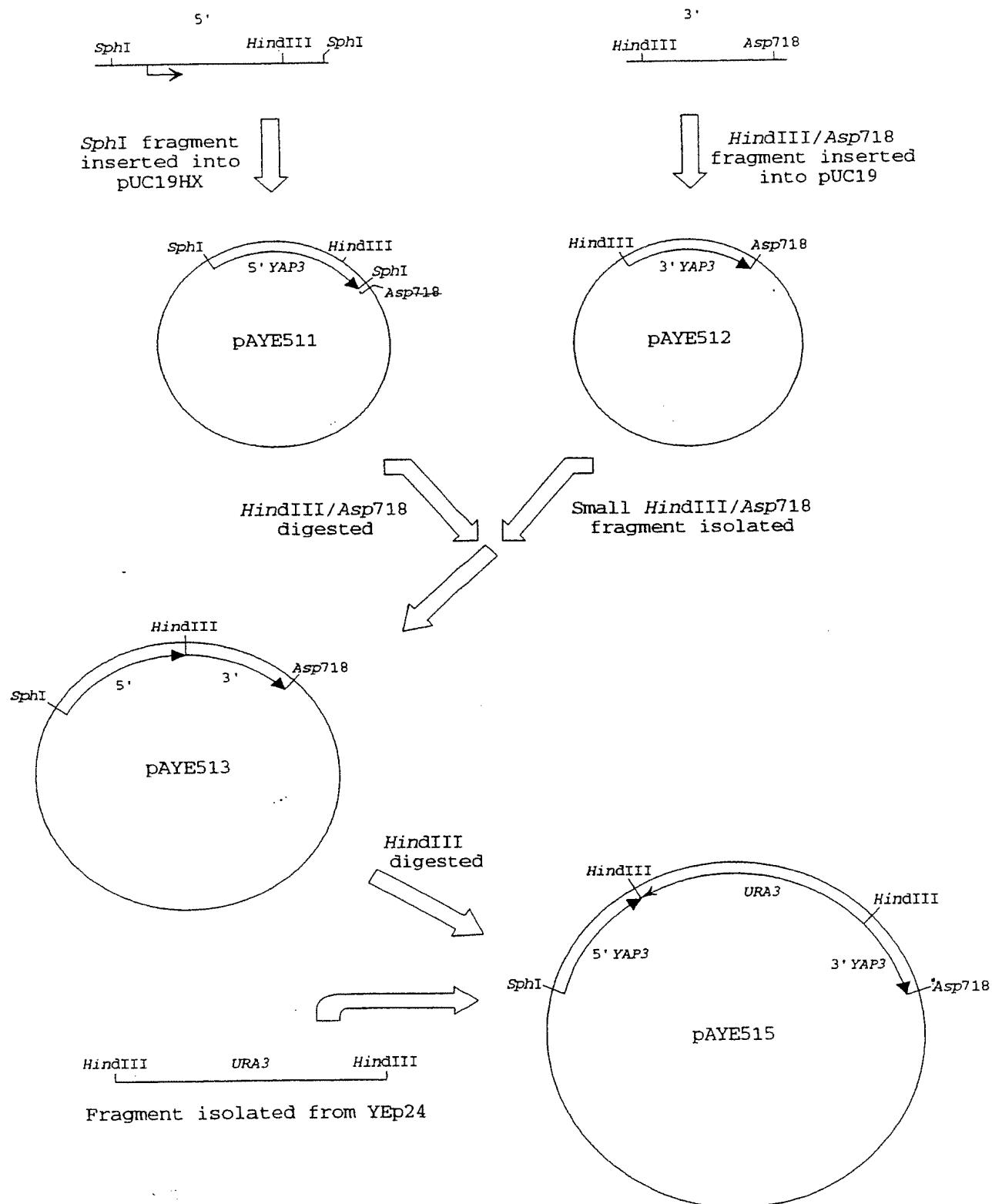


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Figure 2

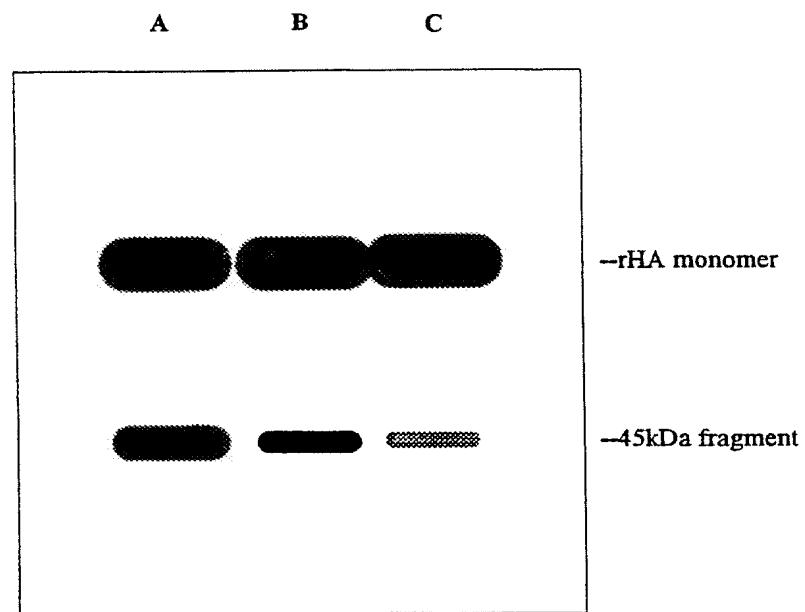
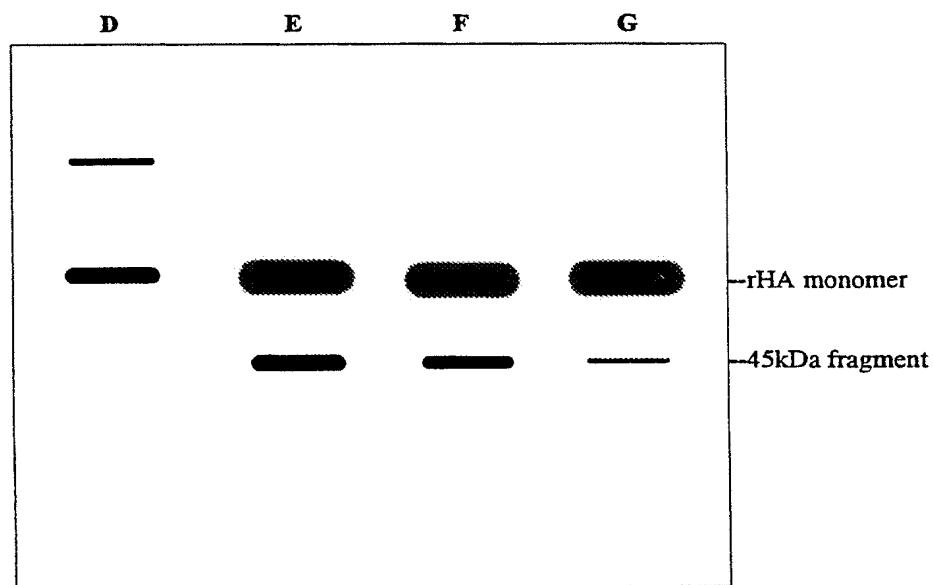
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Figure 3

5' and 3' regions of YAP3 obtained by PCR:

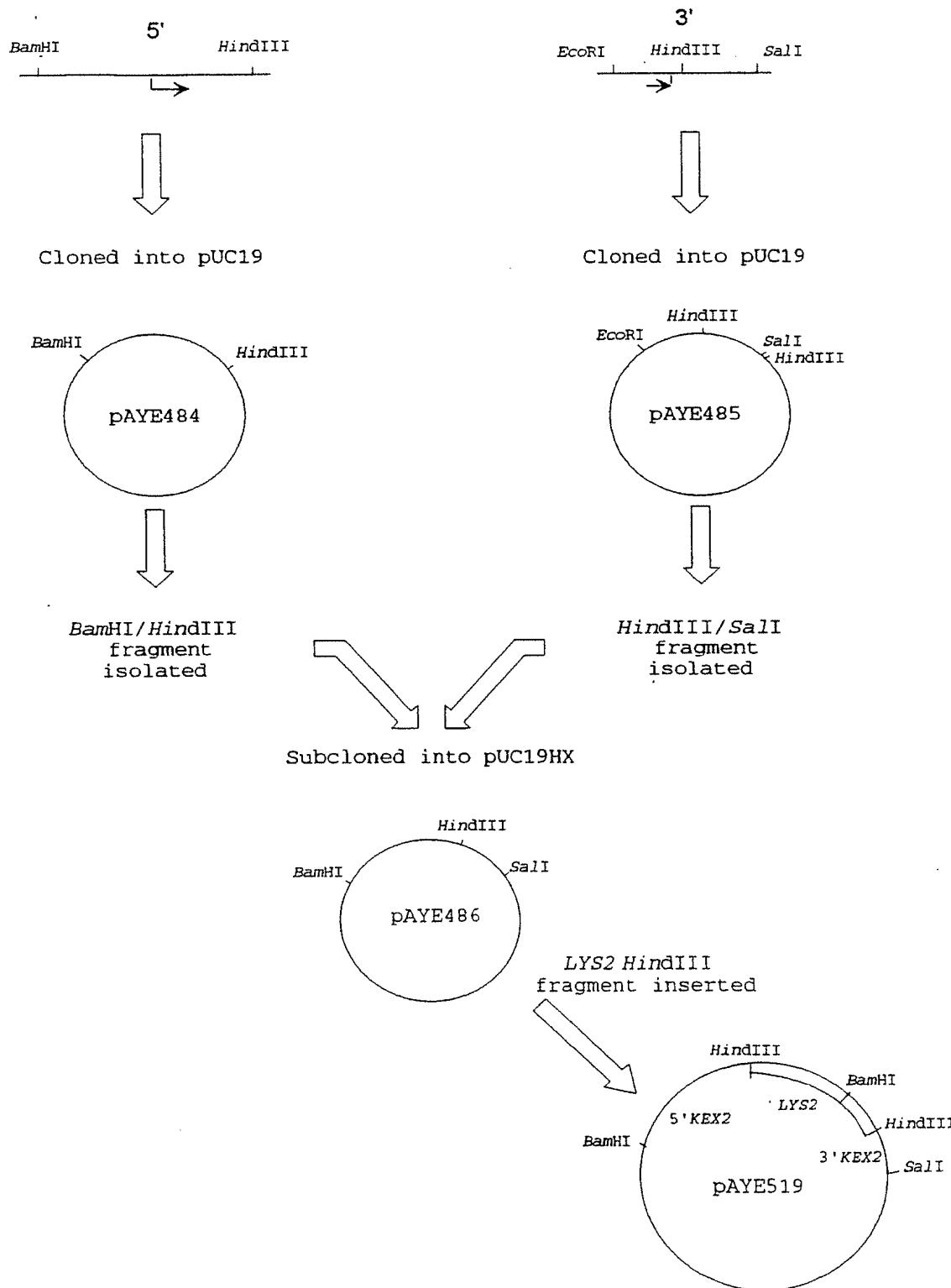


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Figure 4Figure 5

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Figure 6

5' and 3' regions of *KEX2* obtained by PCR:



INTERNATIONAL SEARCH REPORT

International Application No

PCT/GB 95/00434

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 6 C12N15/12 C12N1/21
 C12N1/20

A61K38/38

C12P21/02

C12N15/11

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
 IPC 6 C12N A61K C12P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, A	BIOCHIMIE, vol. 76, 1994 pages 226-233, Y. BOURBONNAIS ET AL. 'Cleavage of prosomatostatins by the yeast Yap3 and Kex2 endoprotease' *see the whole article* ---	1-21
Y	BIOTECHNOLOGY, vol. 8, no. 1, 1990 pages 42-46, D. SLEEP ET AL. 'The secretion of human serum albumin from the yeast S. cerevisiae using five different leader sequences' *see the whole article* ---	1-21 -/-

 Further documents are listed in the continuation of box C. Patent family members are listed in annex.

* Special categories of cited documents :

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- *P* document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

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Date of the actual completion of the international search

Date of mailing of the international search report

23 June 1995

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INTERNATIONAL SEARCH REPORT

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	EMBO JOURNAL, vol. 12, no. 1, 1993 pages 285-294, Y. BOURBONNAIS ET AL. 'Isolation and characterisation of S. cerevisiae mutants defective in somatostatin expression...' *see the whole article* ---	1-21
Y	YEAST, vol. 6, 1990 pages 127-137, M. EGEL-MITANI ET AL. 'A novel aspartyl protease allowing KEX2-independent MF alfa prophero-mone processing in yeast' *see the whole article* -----	1-21